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(54) **Photosensitive resin and method
for manufacture thereof**

(57) A photosensitive resin is
produced by causing a styrylpyridinium
salt possessing a formyl group or an
acetal group to react upon polyvinyl
alcohol.

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SPECIFICATION

Photosensitive resin and method for manufacture thereof

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This invention relates to a photosensitive resin of high solubility in water and high sensitivity and to a method for the manufacture of this resin.

Photosensitive resins have been used as raw materials for printing plates, as photoresists for photo-etching and photomilling operations and as photosensitive vehicles for paints and printing inks. In recent years, studies have been conducted on the use of these resins for the immobilization of

15 enzymes.

The photosensitive resins which have been known are those having azide group, cinnamoyl group, acryloyl group, etc. as their photosensitive residues. As immobilizing carriers for enzymes, however, these conventional photosensitive resins are not considered to be satisfactory because they have serious problems such as the adverse effects they bring about on enzymes and the solubility they exhibit in water.

The screen printing technique involves spreading gauze on a frame, forming on the gauze a membrane having patterns and effecting printing through the patterns.

Ammonium dichromate-polyvinyl alcohol has generally been used as a photosensitive resin solution for use on a screen printing plate. However, this resin solution has disadvantages in that upon application to gauze, dark reaction of the resin proceeds rapidly and in that industrial pollution is caused due to hexavalent chromium.

A photosensitive resin solution having a composition of polyvinyl alcohol di-azo groups has also been used. However, this resin solution has disadvantages in that the membrane formed by applying this resin solution to gauze exhibits lower photosensitivity than that formed by use of the above-mentioned ammonium dichromate-polyvinyl alcohol and in that dark reaction of these di-azo groups proceeds gradually though later than that of the above-mentioned ammonium dichromate.

That is to say, a photosensitive resin solution for screen printing is required (1) to have no metal incorporated therein which can possibly cause industrial pollution from an environmental conservation point of view, (2) to prevent dark reaction of the resin and (3) to form a membrane exhibiting higher photosensitivity than the conventionally known photosensitive agents so as to obtain an advance in operation efficiency.

As a photosensitive high-molecular compound suitable for such new uses, the inventors formerly synthesized polymers possessing a styrylpyridinium group by the reaction of styrylpyridiniums with the polymer of 2-chloroethyl vinyl ether and demonstrated this polymer to be suitable for the photo-immobilization of enzymes [Ichimura and Watanabe: Collection of Manuscripts II for the 37th

Spring Annual Meeting of the Japan Chemical Society, page 1133 (1978) and Chemistry Letters, page 1289 (1978)].

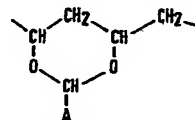
The inventors continued an advanced study with a view to further improving the photosensitivity of this polymer. As a result, they have ascertained that the desired improvement is attained by causing a polyvinyl alcohol to incorporate therein a styrylpyridinium group. The present invention has issued from this knowledge.

An object of this invention is to provide a novel photosensitive resin which possesses high solubility in water and high sensitivity and, therefore, is suitable as a photoresist, a photosensitive vehicle and a screen printing plate and also as an immobilizing carrier for enzymes.

Another object of this invention is to provide a method for the manufacture of the novel photosensitive resin described above.

According to the present invention, there is provided a photosensitive resin comprising a polyvinyl alcohol derivative having incorporated therein a constituent unit of the generic formula:

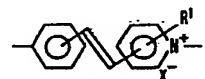
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wherein, A is one member selected from the group consisting of

(A) a group of the generic formula:

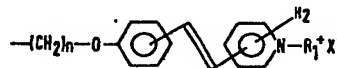
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wherein, R represents one member selected from the class consisting of a hydrogen atom, alkyl groups and lower hydroxyalkyl groups, R' one member selected from the class consisting of a hydrogen atom and alkyl groups and X⁻ a strongly acidic ion, and

(B) a group of the generic formula:

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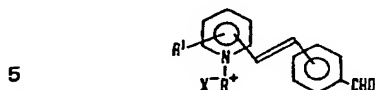


wherein, R₁ represents one member selected from the class consisting of a hydrogen atom, alkyl groups and aralkyl groups, R₂ one member selected from the class consisting of a hydrogen atom and alkyl groups, n an integer having a value between 1 and 6 inclusive and X⁻ a strongly acidic ion.

The invention also provides a method for the manufacture of the above-mentioned photosensitive resin, which method comprises causing one member selected from the group consisting of polyvinyl alcohols and partially saponified polyvinyl acetates to react with one member selected from the group consisting of

(1) styrylpyridinium salts possessing a formyl group

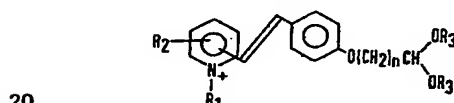
and represented by the generic formula:



wherein, R represents one member selected from the class consisting of a hydrogen atom, alkyl

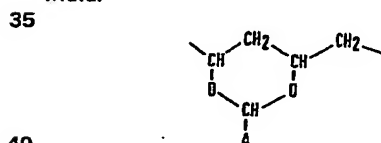
10 groups and lower hydroxyalkyl groups, R' one member selected from the class consisting of a hydrogen atom and alkyl groups and X⁻ a strongly acidic ion, and

15 (2) styrylpyridinium salts possessing an acetal group and represented by the generic formula:

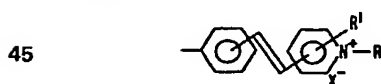


wherein, R1 represents one member selected from the class consisting of a hydrogen atom, alkyl groups and aralkyl groups, R2 one member selected from the class consisting of a hydrogen atom and alkyl groups, two R3's one member selected from the class consisting of an alkyl group and a phenyl group in their respective forms and in a combined form an alkenyl group, n an integer having a value between 1 and 6 inclusive and X⁻ a strongly acidic ion.

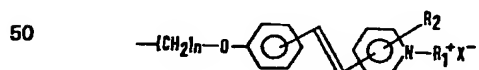
30 The photosensitive resin of the present invention has a chemical structure formed of a polyvinyl alcohol derivative having incorporated therein a photosensitive group of the following generic formula.



40 wherein, A represents either a group of the generic formula

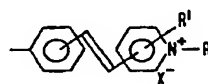


or a group of the generic formula



where, R represents one member selected from the class consisting of a hydrogen atom, alkyl groups and lower hydroxyalkyl groups, R' one member selected from the class consisting of a hydrogen atom and alkyl groups, R1 one member selected from the class consisting of a hydrogen atom, alkyl groups and aralkyl groups, R2 one member selected from the class consisting of a hydrogen atom and alkyl groups, X⁻ a strongly acidic ion and n an integer having a value between 1 and 6 inclusive.

Hereinafter, the generic formula:



70 will be denoted as S and the generic formula:



75 as T.

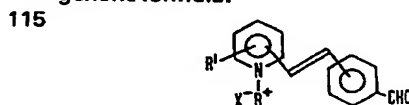
The words "polyvinyl alcohol derivative" as used in the present specification shall mean polyvinyl alcohols and partially saponified polyvinyl acetates and, thus also embraces mixtures containing the polymers of both vinyl alcohol and vinyl acetate.

80 The photosensitive resin of the present invention, therefore, has as its essential constituent units, the aforementioned photosensitive group and the group constituting the aforementioned polyvinyl alcohol derivative. The ratio of these constituent units, specifically the ratio (molar) of the photosensitive group unit to the vinyl alcohol group unit or vinyl acetate group units is desired to fall in the range of from 0.5:99.5 to 10:90. Although the photosensitive group

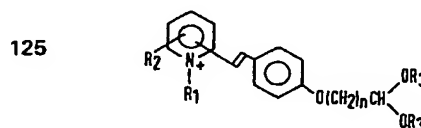
90 in the photosensitive resin possesses hydrophilicity in itself, the resin loses its solubility in water when the photosensitive group unit is present in excess of the upper limit. The proportion of the photosensitive group unit present in the resin, however, can be raised past the upper limit when a polar solvent such as an aqueous alcohol, dimethylformamide, formamide or dimethylsulfoxide or a mixed solvent consisting of such a polar solvent and water as the solvent.

100 The photosensitive resin of the present invention is a novel substance which exhibits high solubility in water and high sensitivity. In an aqueous solution, the photosensitive resin shows the absorption maximum in the ultraviolet spectrum in the neighborhood of 340 nm where the photosensitive group S is used or 360 nm where the photosensitive group T is used. This resin is insolubilized on exposure to a light having a wavelength of up to 460 nm in the former case or up to 500 nm in the latter case.

110 The photosensitive resin of the present invention can be produced by reacting a polyvinyl alcohol or a partially saponified polyvinyl acetate with a styrylpyridinium salt possessing a formyl group of the generic formula:



120 (wherein, R, R' and X⁻ are as defined above) or a styrylpyridinium salt possessing an acetal group of the generic formula:

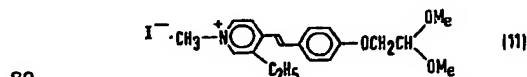
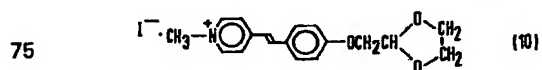
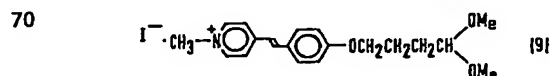
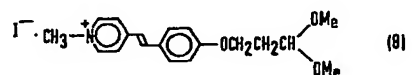
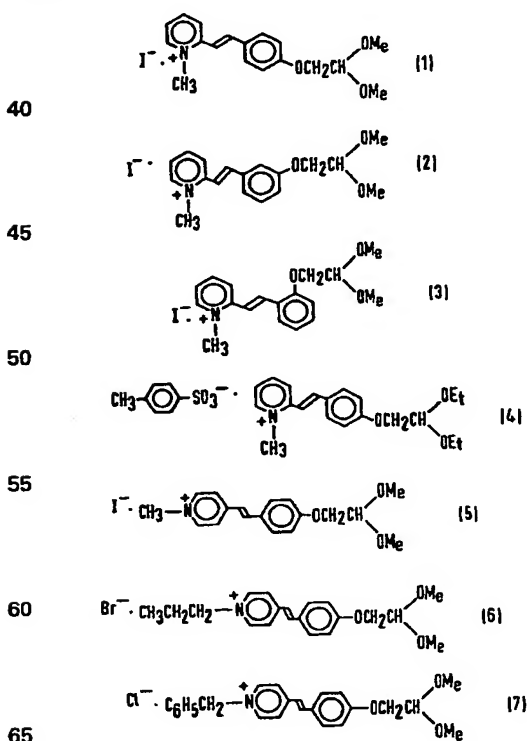


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(wherein, R_1 , R_2 , n and X^- are as defined above and the two R_3 's each represent an alkyl group or a phenyl group or, in a combined form, represent one alkenyl group).

- 5 The styrylpyridinium salt to be used as the raw material for the photosensitive resin of this invention is a novel compound. Examples of such styrylpyridinium salts possessing a formyl group include α - (p - formylstyryl) - pyridinium, γ - (p - formylstyryl) - pyridinium, α - (m - formylstyryl) - pyridinium, N - methyl - α - (p - formylstyryl) - pyridinium, N - methyl - β - (p - formylstyryl) - pyridinium, N - methyl - α - (m - formylstyryl) - pyridinium, N - methyl - α - (o - formylstyryl) - pyridinium, N - ethyl - α - (p - formylstyryl) - pyridinium, N - (2 - hydroxyethyl) - α - (p - formylstyryl) - pyridinium, N - (2 - hydroxyethyl) - γ - (p - formylstyryl) - pyridinium, N - allyl - α - (p - formylstyryl) - pyridinium, N - methyl - γ - (p - formylstyryl) - pyridinium, N - methyl - γ - (m - formylstyryl) - pyridinium, N - benzyl - α - (p - formylstyryl) - pyridinium, N - benzyl - γ - (p - formylstyryl) - pyridinium and N - carbamoylmethyl - γ - (p - formylstyryl) - pyridinium. These quaternary salts may be present in the form of hydrochlorides, hydrobromides, hydroiodides, perchlorates, tetrafluoroborates, methosulfates, phosphates, sulfates, methane-sulfonates and p - toluene - sulfonates. These compounds are obtained by subjecting the corresponding picolines or N - alkyl - picolinium salts to the condensation with aromatic dialdehydes or formyl - benzaldehydes, and they are used in the form of formylstyrylpyridine salts or quaternized salts.

Examples of the styrylpyridinium salts possessing an acetal group include those listed below.



These compounds are obtained by subjecting the corresponding N - alkyl - picolinium salts to the condensation with formyl - phenoxyacetal.

- 85 The polyvinyl alcohol may partly contain an unsaponified acetyl group. This partial inclusion of the group rather meets the purpose of particularly enhancing the solubility of the resin in water. The content of this group is desired not to exceed about 30 mol %. This means that the saponification ratio is desired to be not less than 70%. The degree of polymerization is advantageous in the range of from 400 to 3,000. When the polymerization degree is too small, the time of exposure required for the insolubilization of the resin is notably lengthened. When the polymerization degree is too large, the viscosity is increased even to a point where the actual use of resin is impeded.

- 100 The high-polymer acetalization reaction, i.e. the aforementioned reaction of the polyvinyl alcohol or partially saponified polyvinyl acetate upon the styrylpyridinium salt possessing a formyl group, or the high-polymer interacetalization reaction, i.e. the aforementioned reaction of the polyvinyl alcohol or partially saponified polyvinyl acetate upon the styrylpyridinium salt possessing an acetal group, can be advantageously catalyzed with an acid in water as the medium. In this reaction, the amount of the styrylpyridinium salt to be used is desired to be such that the proportion of this salt relative to the vinyl alcohol unit of the polyvinyl alcohol or partially saponified polyvinyl acetate falls in the range of from 0.3 to 20 mol %. As the catalyst for this reaction, any of inorganic and organic acids may be used.
- 105 Examples of such acids include hydrochloric acid, sulfuric acid, phosphoric acid, perchloric acid, hydrobromofluoric acid, methane - sulfonic acid and p - toluene - sulfonic acid. Since the reaction time decreases with the increasing amount of the acid thus added for catalysis, it is desirable to increase the amount of the acid as much as permissible. For practical purposes, however, the amount of the added acid generally suffices in the range of from 0.01 to 5 normalities. The reaction is carried out at temperatures in the range of from room temperature to 100°C. The reaction time is sufficient in the range of from one to 24 hours in the former reaction and in the range of from one to 48 hours in the latter reaction.
- 120 In this reaction, the optimum polyvinyl alcohol

derivative concentration in the reaction is in the range of from 2 to 20 W/W%.

The progress of the polymer reactions can be traced by precipitating the polymer in alcohol and determining the absorbance due to the styryl-pyridinium group in the neighborhood of 340 nm.

The reaction solutions increase photosensitivity as the reaction proceeds. The progress of this reaction, therefore, can easily be traced measuring the degree of photosensitivity of the reaction solution.

In either case, the solution obtained on completion of the reaction can be used, without purification as a photosensitive liquid of high sensitivity. Desired purification of the resin is accomplished after completion of the reaction by pouring the resultant reaction mixture into a large volume of a non-solvent such as acetone, ethanol or dioxane or a coagulation bath containing such substance as sodium sulfate or potassium sulfate and thereby precipitating the photosensitive resin, separating the precipitated resin from a solvent or a bath and washing the separated resin with alcohol. Thorough removal of the trace of the acid used as the catalyst is attained by washing the resin with alcohol containing a small amount of ammonia or giving the resin repeated reprecipitation.

The photosensitive resin of this invention which contains the photosensitive group S is soluble in water and exhibits high sensitivity. It has a sensitivity equal to or some tens of times as high as that of polyvinyl cinnamate sensitized with 5-nitro-acenaphthene. No other resin has been heretofore known to acquire amply high photosensitivity as the result of the incorporation of a photosensitive group in a very low proportion without use of any additional sensitizer. The resin of the present invention proves to be quite economic because it effectively functions with a very small content of the photosensitive group.

In the case of the resin of this invention which contains the photosensitive group T, the absorption zone expands in the direction of increasing wavelength and comes to exhibit increased sensitivity because the oxy group is directly linked and conjugated with the styrylpyridinium group. This resin has the sensitivity equal to or ten-odd times as high as that of polyvinyl cinnamate sensitized with 5-nitroacenaphthene. The photosensitive group need be present in the resin only in extremely low amounts in the neighborhood of 1 mol % and, the resin is characterized by acquiring amply high photosensitivity without use of any additional sensitizer. The sensitivity of the resin is not affected at all by temperature; virtually no difference is found between that at 80°C and that at -76°C, for example.

By suitable adaptation of the characteristic properties described above, the photosensitive resins of the present invention can be made useful as photoresists, photomilling agents and other printing stocks, as bases for water paints and as vehicles for printing inks. Further, because the points of crosslinking are attained through dimerization of styrylpyridinium groups and, therefore, the crosslinking reaction proceeds in a specific manner in the resin, this resin proves to be highly useful as an immobiliz-

ing carrier for biofunctional materials such as enzymes, sectioned live cells and microorganisms. In view of the adhesive strength inherent in its polyvinyl alcohol unit, the resins also prove to be useful as adhesives and binders of the type allowed to manifest their latent properties in an enhanced manner upon exposure to light.

The photosensitive resin of the present invention can be made useful as a membrane for use in screen printing. The photosensitive resin having a high-molecular emulsion mixed therewith can also be used as the membrane.

Typical examples of high-molecular emulsion include vinylacetate emulsion, acryl emulsion, ethylene-vinylacetate emulsion, ethylene-acryl emulsion, SBR latex, silicone resin emulsion, vinyl chloride emulsion and vinylidene chloride emulsion.

The mixing ratio of the solid phase of the emulsion to the polyvinyl alcohol derivative of the present invention having a photosensitive group is desirably 1:0.05-2. The mixture may generally be composed of 0.01-3 part by weight of the solution having 5-20% by weight of a resin per part of a high-molecular emulsion having 20-70% of solid phase. As occasion demands, a thickener or a small amount of dye may further be mixed with the photosensitive resin of the present invention. The addition of such dye makes it easy to detect the presence of pinholes.

The emulsive, photosensitive compound of the present invention can be preserved for a long period of time and eliminates the disadvantages of the existing method wherein a photosensitive agent and an emulsion are mixed immediately before the mixture is put to practical use since the compound can be used as a photosensitive solution without requiring such mixing treatment.

Now, the present invention will be described more specifically by reference to working examples thereof. Examples 1-6 concern raw materials for the resins of the present invention and Examples 7-26 the resins of the invention.

Example 1:

In a mixed solution consisting of 22.4 g of acetic anhydride and 12.0 g of glacial acetic acid, 18.6 g of α -picoline and 33.5 g of terephthal dialdehyde were dissolved and refluxed under application of heat for eight hours. The resultant reaction solution was cooled, then dissolved in 200 ml of dichloromethane, washed with water and washed again with a diluted aqueous alkali. Subsequently, the reaction product was extracted repeatedly with 1N hydrochloric acid and the extracted product, upon alkalization, immediately gave rise to precipitation of crystals. The crystals were separated by filtration and dried. The crystalline product was dissolved in hot methyl cyclohexane, freed from insoluble material by decantation and recrystallized. Consequently, there were obtained 30.2 g of γ -(p-formylstyryl)-pyridine crystals having a melting point of 84 to 86.5°C. By the same procedure, but using γ -picoline and the dialdehyde, γ -(p-formylstyryl)-pyridine having a melting point of 111 to 113°C was obtained.

Example 2:

In 80 ml of ethyl acetate, 10.0 g of α -(p-formyls-

tyrilyl) - pyridine and 7.2 g of dimethyl sulfate were dissolved and refluxed under application of heat for five hours. Two hours after the completion of the reaction, the reaction solution was cooled. The reaction product was separated by filtration, washed with ethyl acetate and dried. Consequently, there was obtained 15 g of yellow N - methyl - α - (p - formylstyryl) - pyridinium methosulfate having a melting point of 151 to 154°C.

Example 3:

In 20 ml of ethanol were hot dissolved 5.00 g of N - methyl - α - picolinium p - toluenesulfonate and 10.0 g of terephthal dialdehyde. The resultant mixture, after addition thereto of five drops of piperidine, was refluxed for three hours. The resultant reaction solution was cooled, freed from about 10 ml of ethanol through distillation under vacuum, mixed with ethyl acetate to give rise to a yellow precipitate. The precipitate was separated by decantation and washed twice with ethylacetate. When this separated precipitate was dissolved in hot ethanol and gradually mixed with ethyl acetate, there were precipitated bright yellow crystals. The crystals, when separated by filtration, washed with ethyl acetate and dried, produced 5.57 g of N - methyl - α - (p - formylstyryl) - pyridinium p - toluenesulfonate.

Example 4:

In 50 ml of methanol were dissolved 50 g of p - hydroxybenzaldehyde and 23 g of potassium hydroxide. The resultant was freed from the solvent through distillation under vacuum and then dried, to produce a potassium salt. This salt was dissolved in 60 ml of N - methylpyrrolidone, mixed with 70 g of chloroacetaldehyde dimethyl acetal and refluxed at 150°C for 15 hours. The resultant reaction solution was cooled, mixed with 200 ml of dichloromethane, washed three times with water and then washed twice with 50 ml of an aqueous 20 weight % caustic soda solution to extract the unreacted portion of hydroxylaldehyde. The organic layer was washed once with water, dried over anhydrous potassium carbonate and then freed from dichloromethane through distillation under vacuum. When the residual oil was distilled under vacuum, there were obtained N - methylpyrrolidone (bp₃ 135°C) and then p - formylphenoxyacetaldehyde dimethylacetal (bp₃ 145°C). The product obtained through the redistillation totalled 34.2 g.

Example 5:

In 10 ml of 2 - ethoxyethyl alcohol were dissolved 4.88 g of m - hydroxybenzaldehyde and 1.65 g of caustic soda. The resultant solution, after addition thereto of 7.4 g of bromoacetaldehyde dimethylacetal, was refluxed for 20 hours. The resultant reaction solution was cooled, mixed with benzene, washed once with water and then washed with a dilute alkali solution until the hydroxylaldehyde was completely removed. When the benzene solution was desiccated over anhydrous potassium carbonate and then distilled, there was obtained 3.6 g of m - formylphenoxyacetaldehyde dimethyl acetal (bp₃ 138°C). By repeating this procedure faithfully by using o - hydroxybenzaldehyde, there were obtained o - formylphenoxyacetaldehyde dimethylacetal (bp₃ 147°C) and γ - (p - formylphenoxy) - butylaldehyde

dimethyl acetal (bp₃ 168°C).

Example 6:

In 7 ml of methanol were dissolved 1.61 g of N - methyl - α - picolinium iodide and 1.70 g of p - formylphenoxyacetaldehyde dimethyl acetal. The resultant mixture, after addition thereto of 0.3 ml of piperidine, was refluxed for four hours. When the reaction solution was cooled, there ensued precipitation of crystals. The crystals were recovered, washed with a small amount of cooled methanol and washed thoroughly with acetone. Thus, there was obtained 2.11 g of 1 - methyl - 2 - {p - (2,2 - dimethoxyethoxy) - styryl} - pyridinium iodide having a melting point of 192 to 197°C. By following the same procedure, but using the n - methyl - α - picoliniumiodide in combination with m - formylphenoxyacetaldehyde dimethylacetal and n - methyl - γ - picolinium p - toluenesulfonate in combination with p - formylphenoxyacetaldehyde dimethylacetal, as the raw materials, there were respectively obtained 1 - methyl - 2 - {m - (2,2 - dimethoxyethoxy) - styryl} - pyridinium iodide having a melting point of 181 to 186°C, 1 - methyl - 2 - {o - (2,2 - dimethoxyethoxy) - styryl} - pyridinium iodide having a melting point of 169 to 173°C, and 1 - methyl - 4 - {p - (2,2 - dimethoxyethoxy) - styryl} - pyridinium p - toluenesulfonate having a melting point of 219 to 226°C.

Example 7:

In 28 ml of water, 2 g of partially saponified polyvinyl alcohol (87% of saponification ratio) having a polymerization degree of 500 was dissolved. In the resultant aqueous solution was dissolved 800 mg of 1 - methyl - 2 - (p - formylstyryl) - pyridinium p - toluenesulfonate. The yellow solution which was formed consequently was shaken at 60°C for 15 hours in the presence of 1 g of p - toluenesulfonate added thereto. The shaken mixture was introduced dropwise into a large volume of acetone, and the precipitate produced in the acetone was separated by decantation and washed twice with acetone. The washed precipitate was stirred in ethanol containing a small amount of ammonia for 30 minutes, then separated by filtration and dried. Thus, 1.89 g of a polymer was obtained. In its aqueous solution, this polymer showed a maximum absorption at 337 nm. By use of the absorption coefficient of 1 - methyl - 2 - styrylpyridinium p - toluenesulfonate, this polymer was demonstrated to possess 1.80 mol % of styrylpyridinium unit. A film produced from the aqueous solution produced a clear negative image when it was exposed for 30 seconds to the beam from a 450-W high pressure mercury lamp. The sensitivity of the film relative to the sensitivity of the polyvinyl cinnamate having a polymerization degree of 1700 and sensitized with 5 - nitroacenaphthene (10 W/W%) (hereinafter referred to as "relative sensitivity") was 2.5.

Example 8:

In 1 g of an aqueous 10 W/W% solution of 87% saponified polyvinyl alcohol having a polymerization degree of 500, 30 mg of 1 - (2 - hydroxyethyl) - 2 - (p - formylstyryl) - pyridinium chloride was dissolved. The resultant mixture was stirred at 70°C for 12 hours in the presence of 100 mg of p - toluene sulfonic acid added thereto. The resultant reaction solu-

tion was introduced dropwise into a large volume of ethanol. The precipitate consequently formed in the solution was separated by filtration, washed with ethanol and dried, to give 92 mg of a product. The content of styrylpyridinium group introduced in the product was 1.50%. The relative sensitivity of the resin was found to be 1.1.

Example 9:

In 2 g of an aqueous 5 W/W% solution of 87% saponified polyvinyl alcohol having a polymerization degree of 500, 80 mg of 1 - methyl - 2 - (p - formylstyryl) - pyridinium p - toluenesulfonate was dissolved. In the presence of 100 mg of 80% sulfuric acid added thereto, the resultant mixture was stirred at 60°C for 15 hours. The resultant reaction solution was poured into a large volume of ethanol, and the precipitate formed consequently was separated through four times of decantation and washed with ethanol. The separated precipitate was immersed for 30 minutes in ethanol containing a small amount of ammonia. The resin was separated from the bath through filtration, washed with ethanol and dried. Consequently, there was obtained 107 mg of a product having 0.65 mol % of styrylpyridinium unit. The resin's relative sensitivity was found to be 1.3.

Example 10:

To 2 g of an aqueous 5 W/W% solution of 87% saponified polyvinyl alcohol having a polymerization degree of 500, 80 mg of 1 - methyl - 2 - (m - formylstyryl) - pyridinium p - toluenesulfonate was added and the resultant mixture was stirred at 75°C for 15 hours. The resultant reaction solution was poured into a large volume of acetone and then washed through decantation twice with acetone, twice with ethanol and once with ammonia-containing ethanol. Thus, there was obtained 110 mg of a product. By analysis with the ultra-violet absorption spectrometry, the resin was demonstrated to have a styrylpyridinium group content of 1.9 mol %. The film of the resin had a relative sensitivity of 1.5.

Example 11:

In 5 g of an aqueous 5 W/W% solution of thoroughly saponified polyvinyl alcohol having a polymerization degree of 2000, 80 mg of 1 - methyl - 2 - (p - formylstyryl) - pyridinium p - toluenesulfonate was dissolved. In the presence of 200 mg of p - toluene sulfonic acid added thereto, the resultant mixture was stirred at 60°C for 15 hours. The reaction solution obtained consequently was poured into a large volume of acetone, giving rise to a precipitate. This precipitate was dried twice with acetone and once with ethanol and dried. Consequently, 263 mg of a polymer was obtained. The resin was found to have a styrylpyridinium group content of 2.10% and a relative sensitivity of 18.

Example 12:

In 2 g of an aqueous 5 W/W% solution of 87% saponified polyvinyl alcohol having a polymerization degree of 2400, 60 mg of 1 - methyl - 2 - (p - formylstyryl) - pyridinium p - toluenesulfonate was dissolved. The resultant mixture, after addition thereto of 500 mg of p - toluene sulfonic acid, was stirred at 60°C for 15 hours. The resultant reaction solution was introduced dropwise into a large volume of acetone. The precipitate consequently produced was

washed twice with acetone and once with ethanol, then immersed in ammonia-containing ethanol for 30 minutes, separated and dried. Thus, 113 mg of a polymer was obtained. The resin had a styrylpyridinium unit of 2.0 mol%. The film of this resin showed a relative sensitivity of 50.

Example 13:

In 10 ml of water, 200 mg of 87% saponified polyvinyl alcohol having a polymerization degree of 500 was dissolved. The mixture, after addition thereto 40 mg of 1 - methyl - 2 - (p - formylstyryl) - pyridinium iodide and 100 mg of 85% phosphoric acid, was stirred at 60°C for three hours. The reaction solution was immediately poured into a large volume of acetone. The precipitate consequently formed was washed thoroughly three times with methanol and then dried. Thus, 205 mg of a polymer was produced. This resin was found to have a styrylpyridinium unit of 1.7 mol% and a relative sensitivity of 2.0.

Example 14:

In 2 ml of water, 400 mg of 87% saponified polyvinyl alcohol having a polymerization degree of 500 was dissolved. The mixture, after addition thereto of 40 mg of 1 - methyl - 4 - (p - formylstyryl) - pyridinium p - toluenesulfonate and 0.5 g of 85% phosphoric acid, was stirred at 65°C for ten hours. The reaction solution was poured into a large volume of acetone. The precipitate consequently produced was washed four times with methanol and then vacuum dried. Thus, 399 mg of a polymer was obtained. This resin was found to have a styrylpyridinium group unit of 0.70 mol% and a relative sensitivity of 0.7.

Example 15:

In 15 ml of water, 1 g of 87% saponified polyvinyl alcohol having a polymerization degree of 1700 was dissolved. The mixture, after addition thereto of 100 mg of 1 - methyl - 4 - (p - formylstyryl) - pyridinium p - toluenesulfonate and 0.5 g of 85% phosphoric acid, was stirred at 80°C for eight hours. The reaction solution was poured into a large volume of acetone to give rise to a precipitate. The precipitate was thoroughly disintegrated and washed three times with methanol. After it was confirmed that the washings no longer assumed a yellow color, the product was vacuum dried. Thus, 0.95 g of a polymer was obtained. Through absorption spectrometry, this resin was shown to possess the maximum absorption at 343 nm and have a styrylpyridinium group content of 1.02%. The resin showed a relative sensitivity of 12. The film prepared from the reaction solution prior to purification showed a relative sensitivity of 7.

Example 16:

To 10 g of an aqueous 5 W/W% solution of 87% saponified polyvinyl alcohol having a polymerization degree of 1700, 100 mg of 1 - methyl - 2 - (p - formylstyryl) - pyridinium - methosulfate was added. The resultant mixture was thoroughly stirred and mixed with 3 g of 85% phosphoric acid under stirring. Consequently, there was obtained a yellow homogeneous solution. This solution was maintained overnight at 67°C and then poured into a large volume of acetone. The precipitate consequently formed in the

solution was collected. This precipitate was washed twice with ethanol and dissolved in water and reprecipitated in acetone for the purpose of purification. Thus, 0.47 g of a resin was obtained. This resin had a styrylpyridinium unit of 1.20 mol% and a relative sensitivity of 3.0.

Example 17:

To a 10 g of an aqueous 5 W/W% solution of 87% saponified polyvinyl alcohol having a polymerization degree of 500, 30 mg of 1 - methyl - 2 - (p - formylstyryl) - pyridinium - methosulfate was added. The mixture was thoroughly stirred and then mixed with 0.5 ml of 6N hydrochloric acid under stirring, to produce a yellow homogeneous solution. This solution was maintained at 70°C for eight hours. The hot solution was poured into a large volume of acetone. The precipitate consequently formed in the solution was washed three times with methanol and then dried. Thus, 482 mg of a resin was obtained. This resin had a styrylpyridinium unit of 0.45 mol% and showed a relative sensitivity of 0.8.

Example 18:

In 10 g of an aqueous 5 W/W% solution of 87% saponified polyvinyl alcohol having a polymerization degree of 1700, 30 mg of 1 - carbamoylmethyl - 4 - (p - formylstyryl) - pyridinium chloride was dissolved. The mixture, after addition thereto of 1 g of 85% phosphoric acid, was stirred at 50°C for ten hours. The resultant reaction solution was poured into a large volume of acetone. The precipitate consequently formed therein was washed four times with methanol and vacuum dried. Thus, 0.45 g of a resin was obtained. This resin had a styrylpyridinium unit of 0.52 mol% and showed a relative sensitivity of 4.0.

Example 19:

In 210 ml of water, 21 g of 87% saponified polyvinyl alcohol having a polymerization degree of 1700 was dissolved by application heat. The solution was homogeneously mixed with 20 ml of water having dissolved in advance therein 2.1 g of 2 - (p - formylstyryl) - pyridine and 3 g of 85% phosphoric acid. The solution was stirred at 70°C for six hours. The reaction solution had a styrylpyridinium unit of 1.26 mol% based on the polyvinyl alcohol. When this reaction solution was applied in its unmodified form to an aluminum plate and the resultant film was exposed to light and then developed with a weakly acidic aqueous solution, there was obtained a clear negative image. The film's relative sensitivity was found to be 6.7.

Example 20:

In 7 ml of water, 500 mg of 87% saponified polyvinyl acetate having a polymerization degree of 1700 was added. Then, 187 mg of 1 - methyl - 4 - (p - (2,2 - dimethoxyethoxy) - styryl) - pyridinium - methosulfate (see Example 6). The resultant solution, after addition thereto of 0.5 ml of 85% phosphoric acid, was stirred at 60°C for 15 hours. The reaction solution showed a high degree of sensitivity in itself. It was further processed as follows to afford a resin. The yellow reaction solution was poured into a large volume of acetone to produce a resin in the form precipitate. The precipitate was thoroughly washed twice with methanol and vacuum dried. Thus, there was obtained 420 mg of resin. The resin,

in its aqueous solution, showed maximum absorption at 370 nm. The resin was estimated to have a styrylpyridinium unit of 2.20 mol% on the basis of the ultraviolet absorption spectrum. This resin was converted into an aqueous solution, applied to an aluminum plate and dried, and the resultant film was exposed through a negative image to the beam from a 450-W high pressure mercury lamp and then developed with water. Consequently there was obtained a clear image. The resin's relative sensitivity was found to be 30.

When the procedure described above was repeated by using the partially saponified polyvinyl acetate and the phosphoric acid in the same amounts as above, except that the amount of the aforementioned pyridinium salt was changed to 97.1 mg in one test run and 49.5 mg in another test run, there were obtained photosensitive polyvinyl alcohol resins having 1.10 mol% and 0.71 mol% respectively of styrylpyridinium unit. These resins were found to have 12 and 9 respectively of relative sensitivity.

Example 21:

In 6 g of an aqueous 8 weight % solution of 87% saponified polyvinyl acetate (having a polymerization degree of 1700), 63 mg of 1 - methyl - 2 - {o - (2,2 - dimethoxyethoxy) - styryl} - pyridinium iodide was added. The resultant mixture, after addition thereto of 150 mg of p - toluene sulfonic acid, was treated at 60°C for 15 hours to induce a reaction. The yellow reaction solution consequently produced was poured into a large volume of acetone. The resin which occurred in the form of precipitate was washed twice with methanol and then dried under vacuum. Thus, 409 mg of purified resin was obtained. The aqueous solution of this resin showed the maximum absorption at 356 nm. On the basis of the absorption coefficient of the pyridinium salt used as the raw material, the resin was estimated to contain 1.43 mol% of styrylpyridinium group. When tested by the gray scale method, this resin showed a relative sensitivity of 2.7. When the procedure described above was repeated by using 33 mg of the pyridinium salt, there was obtained a resin having a styrylpyridinium unit of 0.81 mol%. The relative sensitivity of this resin was 0.6.

Example 22:

In 6 g of an aqueous 8 weight% solution of 87% saponified polyvinyl acetate (having a polymerization degree of 1700), 60.5 mg of 1 - methyl - 2 - {m - (2,2 - dimethoxyethoxy) - styryl} - pyridinium iodide was homogeneously dissolved. The solution, after addition thereto of 150 mg of p - toluene sulfonic acid, was stirred at 60°C for 15 hours. The reaction solution was poured into a large volume of acetone. The resin consequently produced in the solution was washed twice with methanol and then dried under vacuum. Thus, there was obtained 420 mg of a purified resin. The aqueous solution of this resin showed the maximum absorption at 338 nm. On the basis of the absorption spectrum, the resin was estimated to have a styrylpyridinium content of 1.43%. The resin's relative sensitivity was 7.0. When the procedure described above was repeated by using 32.8 mg of the aforementioned pyridinium salt, there was obtained a resin having a styrylpyridinium unit

of 0.82 mol%. The relative sensitivity of this resin was 2.4.

Example 23:

- In 6 g of an aqueous 8 weight% solution of 87% saponified polyvinyl acetate (having a polymerization degree of 1700), 61.4 mg of 1-methyl-2-{p-(2,2-dimethoxyethoxy)-styryl}-pyridinium iodide was dissolved. The solution was mixed with 150 mg of p-toluene sulfonic acid to produce a homogeneous solution, which was treated at 60°C for 15 hours to induce a reaction. The yellow reaction solution thus formed was poured into a large volume of acetone. The resin which occurred consequently was washed twice with methanol and then dried under vacuum. Thus there was obtained 420 mg of a purified resin. The aqueous solution of this resin showed the maximum absorption at 361 nm. On the basis of the absorption spectrum, this resin was estimated to have a styrylpyridinium unit of 1.27 mol%. This resin had a relative sensitivity of 9.0. When the procedure described above was repeated by using 33.7 mg of the aforementioned pyridinium salt, there was obtained a resin having a styrylpyridinium unit of 0.75 mol%. It showed a relative sensitivity of 4.0.

Example 24:

- In 6 g of an aqueous 8 weight% solution of 87% saponified polyvinyl acetate (having a polymerization degree of 1700), 65.0 mg of 1-methyl-2-{p-(4,4-dimethoxybutoxy)-styryl}-pyridinium iodide obtained from N-methyl- α -picolinium iodide and γ -(p-formylphenoxy)-butylaldehyde dimethylacetal by the procedure of Example 6 was homogeneously dissolved. The resultant solution, after addition thereto of 150 mg of p-toluene sulfonic acid, was treated at 60°C for 15 hours to induce a reaction. The yellow reaction solution was poured into a large volume of acetone. The resin consequently formed was washed twice with methanol and then dried under vacuum. Thus, there was obtained 430 mg of a purified resin. This resin, in its aqueous solution, showed the maximum absorption at 361 nm. On the basis of the absorption spectrum, the resin was estimated to have a styrylpyridinium unit of 1.05 mol%. The resin's relative sensitivity was 3.2.

Example 25:

- 10 g of N-methyl- γ -(p-formylstyryl)-pyridinium methosulfate was dissolved in a solution produced by dissolving in 1150 ml of water 100 g of 87% saponified polyvinyl acetate having a polymerization degree of 2000 and then mixed with 5 g of 85% phosphoric acid. The resultant mixture was stirred at 70°C for 5 hours. To 12 g of the resultant reaction solution, 7.5 g of vinylacetate emulsion (Konishi Shoten: Bond-CH-18) was added and fully mixed. The solution thus obtained was applied to gauze of 150 mesh made of Tetlon and dried, with the result that a membrane of uniform thickness was formed. The membrane was placed at a position about 50 cm distant from a 450W mercury lamp and irradiated through a negative for two minutes, and thereafter vigorously washed with city water. As a result, a clear image appeared on the membrane. This membrane exhibited about five times the photosensitivity

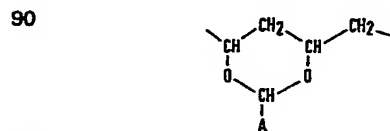
of a membrane formed by use of a commercially available photosensitive solution of di-azo resin-polyvinyl alcohol.

Example 26:

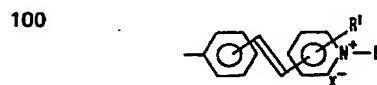
- A mixed solution of N-methyl- γ -(p-formylstyryl)-pyridinium methosulfate, polyvinyl acetate and phosphoric acid was prepared in the same manner as in Example 25. 1.5 parts by weight of silicone resin emulsion (30% solid phase) per part of this solution was well mixed with this solution. The resultant mixture was applied to gauze of 150 mesh made of Tetlon and dried to obtain a membrane. This membrane was irradiated with a super high-pressure mercury lamp and developed with city water. As a result, a clear image appeared on the membrane. This membrane exhibited high strength, high repellence and about ten times the photosensitivity of a membrane formed by use of a commercially available di-azo resin photosensitive agent.

CLAIMS

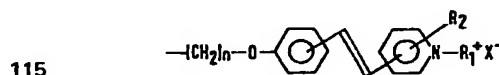
1. A photosensitive resin comprising a polyvinyl alcohol derivative having incorporated therein a constituent unit of the generic formula:



- 95 wherein, A is one member selected from the group consisting of
(A) a group of the generic formula:



- 105 wherein, R represents one member selected from the class consisting of a hydrogen atom, alkyl groups and lower hydroxyalkyl groups, R' one member selected from the class consisting of a hydrogen atom and alkyl groups and X⁻ a strongly acidic ion, and
110 (B) a group of the generic formula:



- wherein, R₁ represents one member selected from the class consisting of a hydrogen atom, alkyl groups and aralkyl groups, R₂ one member selected from the class consisting of a hydrogen atom and alkyl groups, n an integer having a value between 1 and 6 inclusive and X⁻ a strongly acidic ion.

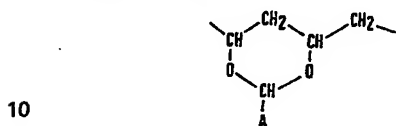
2. The photosensitive resin according to Claim 1, wherein X⁻ represents one member selected from the group consisting of halogen ions, sulfate ion, phosphate ion and p-toluene sulfonate ion.

3. The photosensitive resin according to Claim 1, wherein the polymerization degree of the polyvinyl alcohol falls in the range of from 400 to 3000.

4. The photosensitive resin according to Claim 1,

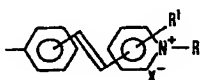
which resin is useful for a screen printing plate.

5. A method for the manufacture of a photosensitive resin comprising a polyvinyl alcohol derivative having incorporated therein a constituent unit of the generic formula:



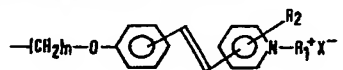
wherein, A is one member selected from the group consisting of
(A) a group of the generic formula:

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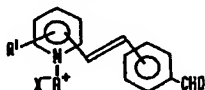
- 20 wherein, R represents one member selected from the class consisting of a hydrogen atom, alkyl groups and lower hydroxyalkyl groups, R' one member selected from the class consisting of a hydrogen and alkyl groups and X⁻ a strongly acidic ion, and
25 (B) a group of the generic formula:

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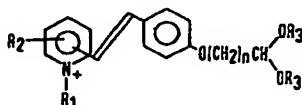
- wherein, R, represents one member selected from the class consisting of a hydrogen atom, alkyl groups and aralkyl groups, R₂ one member selected from the class consisting of a hydrogen atom and alkyl groups, n an integer having a value between 1 and 6 inclusive and X⁻ a strongly acidic ion, which method comprises causing one member selected from the group consisting of polyvinyl alcohols and partially saponified polyvinyl acetates to react with one member selected from the group consisting of
(1) styrylpyridinium salts possessing a formyl group and represented by the generic formula:

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- wherein, R represents one member selected from the class consisting of a hydrogen atom, alkyl groups and lower hydroxyalkyl groups, R' one member selected from the class consisting of a hydrogen and alkyl groups and X⁻ a strongly acidic ion, and
55 (2) styrylpyridinium salts possessing an acetal group and represented by the generic formula:

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- wherein, R₁ represents one member selected from the class consisting of a hydrogen atom, alkyl groups and aralkyl groups, R₂ one member selected from the class consisting of a hydrogen atom and
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alkyl groups, two R₃'s one member selected from the class consisting of an alkyl group and a phenyl group in their respective forms and in a combined form an alkenyl group, n an integer having a value between 1 and 6 inclusive and X⁻ a strongly acidic ion.

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6. The method according to Claim 5, wherein X⁻ represents one member selected from the group consisting of halogen ions, sulfate ion, phosphate ion and p-toluene sulfonate ion.

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7. The method according to Claim 5, wherein the styrylpyridinium salt possessing a formyl group is used in a proportion of from 0.3 to 20 mol% based on one vinyl alcohol unit selected from the group consisting of polyvinyl alcohols and partially saponified polyvinyl acetates.

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8. The method according to Claim 5, wherein the styrylpyridinium salt possessing an acetal group is used in a proportion of from 0.3 to 20 mol% based on one vinyl alcohol unit selected from the group consisting of polyvinyl alcohols and partially saponified polyvinyl acetates.

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9. A photosensitive resin substantially as hereinbefore described.

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10. A method of manufacturing a photosensitive resin substantially as hereinbefore described.

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